

K832827 CORPULS 300Nov 3, 1983
73 days to decisionK832827 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k832827/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 22, 1983
Decision date	Nov 3, 1983
Days to decision	73 days
Third-party review	No

APPLICANT

Company	Automated Medical Products, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1983

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k832827/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 8, 2026